

Photon recycling white light emitting diode based on InGaN multiple quantum well heterostructure

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A numerical method based on the transfer matrix technique is developed to calculate the luminescence spectra of complex layered structures with photon recycling. Using this method we show a strong dependence of the emission spectra on the optical eigenmode structure of the device. The enhancement of the photon recycling and the LED external efficiency can be achieved by placing the active regions inside single or coupled microcavities.

I. INTRODUCTION

Recently, great interest has been shown in the use of light emitting diodes (LEDs) as a light source for illumination [1]. LEDs offer many potential advantages compared to conventional light sources due to their relatively low energy consumption, long lifetime and high shock resistance.

Currently white-light LEDs use photo-excitation of phosphors to convert the blue light from an InGaN/GaN LED into white light. However, phosphor has a broad emission spectrum, thus, white LEDs based on this principle do not have the maximum possible luminous efficacy.

The aim of this work is to investigate the feasibility and limitations of creating a white LED by integration within the same structure of several semiconductor layers emitting three basic colours. The recent progress in the growth of InGaN-based double heterostructures and quantum wells makes this alloy an ideal material for the LED active regions, due to the wide variety of the energy gaps in the InGaN system, covering frequencies from red to ultraviolet. The working regime of such a device can be achieved by electrical pumping of active layers with the widest bandgap (blue regions), and by making use of re-emission of the light, absorbed by all the active regions (so called photon recycling).

II. MODELLING OF PHOTON RECYCLING

We start by writing an expression for the intensity of spontaneous emission from a quantum well (QW) into a bulk dielectric material. The number of photons with energy within the interval $[\hbar\omega, \hbar\omega + d(\hbar\omega)]$, which are emitted from the surface dS into the solid angle $d\Omega_0$ during the time interval dt is given by

$$dN = W_0 dt d(\hbar\omega) dS d\Omega_0 = \frac{\epsilon\omega^3 \hbar e^2 p_{cv}^2 g_{2D}}{2\pi^2 c^3 m_0^2} \left[\int_0^{+\infty} \frac{f_h f_e \Gamma d\varepsilon}{(E^{2D} + \varepsilon)^2 ((E^{2D} + \varepsilon - \hbar\omega)^2 + \Gamma^2)} \right] dt d(\hbar\omega) dS d\Omega_0. \quad (1)$$

Here E^{2D} is the energy gap between the electron and hole quantized levels in the QW, g_{2D} is the reduced two-dimensional density of states, f_e and f_h are the electron and hole occupation probabilities, ϵ is the dielectric constant of the material containing the well, m_0 is the free electron mass, and Γ accounts for the interband relaxation and other broadening mechanisms. The squared momentum matrix element p_{cv}^2 is given in the effective mass approximation for deep QWs by [2]

$$p_{cv}^2 = \frac{p_0^2}{2} (1 + \gamma)$$

for TE modes, and by

$$p_{cv}^2 = \frac{p_0^2}{2} [(1 + \gamma) \cos \theta + (1 - \gamma) \sin \theta]$$

for TM modes. Here θ is the angle between the plane-wave propagation direction and Z -axis (normal to the QW plane), and $\gamma = (E_{2D} - E_g)/(\hbar\omega - E_g)$, where E_g is the bandgap energy of the QW material, and p_0 is the interband momentum matrix element.

We will use the transfer matrix technique in the plane waves basis [3] to analyse the optical properties of layered structures. The plain-wave mode interaction with the QW is described by a QW transfer matrix. If the quantum well width is much smaller than the light wavelength, the transfer matrix has the form:

$$\hat{M} = \begin{pmatrix} 1+Y & Y \\ -Y & 1-Y \end{pmatrix}.$$

Here Y is defined by the two-dimensional QW optical susceptibility χ_{2D} :

$$Y = i \frac{2\pi k_0^2}{k_z} \chi_{2D}, \quad (2)$$

where \mathbf{k} is the wavevector of light, $k_0 = \omega/c$, and χ_{2D} for a single-subband QW is given by [4]

$$\chi_{2D} = \frac{\hbar^2 e^2 p_{cv}^2 g_{2D}}{m_0^2} \int_0^{+\infty} \frac{1 - f_h - f_e}{(E^{2D} + \varepsilon)^2 (E^{2D} + \varepsilon - \hbar\omega - i\Gamma)} d\varepsilon$$

To calculate the rate of photon extraction from a complex structure we consider the interference of the all possible processes, resulting in light emission out of the structure. These calculations require a knowledge of the amplitude transmission and reflection coefficients t_l and r_l for the structure part on the left of the QW, and similar coefficients t_r and r_r for the structure part on the right of the QW. We also need to know the transmission and reflection coefficients t_{QW} and r_{QW} for the quantum well itself. Each of these coefficients can be obtained from the corresponding transfer matrix.

Let us derive, for example, the power, emitted from the right side of the structure. A photon emitted inside the structure to the right can be transmitted directly to the outside medium, or it can be consecutively reflected from the right and left sides of the structure and finally will be transmitted outside and so on. The outgoing electric field, resulting from all these processes is given by the sum:

$$E_{r \rightarrow r} = t_r + r_r r_l^* t_r + r_r r_l^* r_r r_l^* t_r + \dots = \frac{t_r}{1 - r_l^* r_r},$$

where the star in r_l^* indicates, that this coefficient includes the reflection from the emitting QW.

Similarly, photons emitted inside the structure to the left can undergo multiple reflections and eventually escape from the structure to the right. These processes give a second part of the external field:

$$E_{l \rightarrow r} = \frac{r_l t_{QW} t_r}{(1 - r_l^* r_r)(1 - r_{QW} r_l)}$$

Thus, we obtain the expression for the emission from the right side of the structure:

$$dN_{Er} = W_0 * \left| \frac{t_r}{1 - r_l^* r_r} \left[1 + \frac{r_l t_{QW}}{1 - r_{QW} r_l} \right] \right|^2 \frac{\sqrt{\epsilon_r} k_{zr}}{\sqrt{\epsilon_0} k_{z0}} dt d(\hbar\omega) dS d\Omega_e, \quad (3)$$

where the ratio $\sqrt{\epsilon_r} k_{zr} / \sqrt{\epsilon_0} k_{z0}$ accounts for the change in solid angle that is due to refraction for plane waves. The indices r and 0 are related to the outside medium and the layer containing the QW, respectively. To obtain the total density of the external light intensity we have to sum over all QWs and integrate Eq.(3) over the external solid angle Ω_e .

If we neglect the reflection from the QW by substituting $t_{QW} = 1$ and $r_{QW} = 0$ into Eq.(3), the formula for extraction becomes analogous to one obtained using the source-term method [5].

For the quantitative description of the recycling process, we have to calculate the rate of absorption by the quantum well QW_a of the spontaneous emission from the other well, QW_e . Thus, we need to know the induced electric field at the position of QW_a . The power flux balance shows, that the rate of absorption of emitted photons by the unit area of surface of QW_a is given by:

$$W_a = -\frac{1}{2} \sqrt{\epsilon} k_z |E|^2 \text{Re}(Y_{QW_a}) W_{0QW_e}, \quad (4)$$

where E is the complex amplitude of the field at the QW_a location, Y_{QW_a} is defined by Eq.(2), and W_{0QW_e} is defined by Eq.(1). The induced field can be calculated in a similar fashion as it was done for extraction:

$$E = \frac{t(1 + r_{al}^*)}{(1 - r_{er}^* r_{er}^*)(1 - r_{al}^* r_{ar})} \left(1 + \frac{t_{QW_e} r_{er}}{1 - r_{er} r_{QW_e}} \right),$$

where r_{al} (r_{ar}) is a reflection coefficient for the part of structure to the left (right) of the well QW_a , r_{er} is a reflection coefficient for all the layers to the right of the well QW_e , and the coefficients r and t correspond to the part of the structure between QW_e and QW_a . Here we assume that QW_a is on the left of QW_e . The formula for the opposite case is similar.

One of the channels for the photon to escape from the recycling process is to be absorbed in a metallic mirror. Placing such a mirror onto the left side of the structure, and denoting the reflection coefficient of the left part of the structure, excluding the mirror, as r_l , the reflection coefficient from the mirror as r_m , the reflection coefficient for the wave incident from the mirror as r_s , and the transmission coefficient from the QW to the mirror as t , we obtain the following expression for the number of absorbed photons:

$$dN_m = W_0 (1 - |r_m|^2) \left| \frac{t}{(1 - r_l r_r^*)(1 - r_m r_s)} \left(1 + \frac{r_r t_{\text{QW}}}{1 - r_r r_{\text{QW}}} \right) \right|^2 dt d(\hbar\omega) dS d\Omega_0. \quad (5)$$

We restrict our consideration by relatively low pumping levels, which are typical for the diode operation regime. Thus, the expression for the electron density [4] in the one-subband QW can be simplified (temperature is measured in energy units):

$$n = \frac{m_c}{\hbar^2 \pi} T \ln[1 + \exp(\mu_e/T)] \approx \frac{m_c}{\hbar^2 \pi} T \exp(\mu_e/T),$$

and the occupation probabilities can be expressed as:

$$f_e = \frac{1}{1 + \exp(\frac{\varepsilon_e - \mu_e}{T})} \approx \exp\left(\frac{\mu_e - \varepsilon_e}{T}\right) \approx n \frac{\hbar^2 \pi}{m_c T} \exp(-\varepsilon_e/T),$$

$$f_h = \frac{1}{1 + \exp(\frac{\varepsilon_h - \mu_h}{T})} \approx \exp\left(\frac{\mu_h - \varepsilon_h}{T}\right) \approx p \frac{\hbar^2 \pi}{m_{hh} T} \exp(-\varepsilon_h/T),$$

where m_e (m_{hh}) and μ_e (μ_h) are the electron (heavy hole) effective mass and quasi-Fermi-level, respectively. Under the low-pumping assumption we can rewrite W_0 in the form:

$$W_0 = \frac{\epsilon \omega^3 \hbar e^2 p_{cv}^2 g_{2D}}{2\pi^2 c^3 m_0^2} \left[\int_0^{+\infty} \frac{\Gamma \exp(-\varepsilon/T) d\varepsilon}{(E^{2D} + \varepsilon)^2 ((E^{2D} + \varepsilon - \hbar\omega)^2 + \Gamma^2)} \right] np. \quad (6)$$

The total rate of light emission into the external medium can be obtained by substituting Eq.(6) in Eq.(1) and integrating Eq.(1) over the external solid angle and photon energies. As a result the rate of radiative recombination depends on the carrier densities as $R_{ext} = E np$.

Due to the large photon energy separation between different colours, the absorption rate for the short-wavelength light is independent of the carrier density in the narrow-gap QW. For the light emitted from QW_e the rate of absorption in QW_a is given by the expression $R_{abs} = A n_e p_e$, where A is calculated by integrating Eq.(4) over the total solid angle of emission and over the photon energies. Here we assume that all QWs are embedded in layers of dielectric material with the refractive index equal to the highest one in the real structure. This trick allows us to handle the interaction of active layer with the evanescent wave. However, introducing several sufficiently thin layers with high dielectric constant does not alter the optical properties of the structure.

A similar relation holds for the rate of absorption in a metallic mirror: $R_m = M np$. Coefficient M is obtained by substituting Eq.(6) in Eq.(5) and integrating the result over the solid angle and photon energies. Note, that due to charge-neutrality the electron and hole densities in each QW are equal to each other, $n = p$.

The steady-state carrier densities are given by the balance between generation of electron-hole pairs in the QWs and their recombination, both radiative and non-radiative. The generation processes include electric current pumping of the blue QWs and the re-absorption of emitted light throughout the structure. The recombination output goes to external emission and internal losses, which we treat as absorption in the other QWs and non-radiative recombination in the given QW. In our approach this leads to the following equation for each QW:

$$R_i = E_i n_i^2 + A_i n_i^2 + M_i n_i^2 + n_i / \tau_i,$$

where R_i is the pumping rate and τ_i is the non-radiative recombination time. Assuming that τ_i does not depend on the carrier density, we get:

$$n_i = \frac{\sqrt{(1/\tau_i)^2 + 4(A_i + E_i + M_i)R_i} - 1/\tau_i}{2(A_i + E_i + M_i)}. \quad (7)$$

Then the external efficiency for each QW can be obtained as

$$\eta_i = \frac{E_i n_i^2}{(A_i + E_i + M_i)n_i^2 + n_i/\tau_i}.$$

III. RESULTS AND DISCUSSION

We used the numerical method described above to investigate a number of different types of structures. It was revealed, that the emission spectra and external efficiencies depend drastically on the active layer positions and on the mode structure of the device. Figure 1 shows the calculated spontaneous emission for a structure, which represents GaN microcavity containing three QWs and covered on one side by a metallic mirror. The width of the GaN layer and the energies of interband transitions in red, green and blue QWs are chosen in such a way that the structure can be regarded as the $5\lambda/2$ resonator for the normally propagating red-light waves, $6\lambda/2$ resonator for the green-light waves and $7\lambda/2$ resonator for the blue-light waves. We placed each QW in an antinode of a resonant mode, associated with the QW colour. Our calculations were performed for different times of non-radiative recombination ranging from 1 ns to 10 ns, and we assumed the carrier density in blue QW to remain constant. One can see, that if the internal losses are high, the recycling efficiency is low and a blue light only is emitted. When the internal efficiency increases, the emission from the optically pumped QWs becomes comparable with the blue-light intensity. The green-light intensity usually remains smaller than both blue and red because of the strong re-absorption in the red active region. However, this is beneficial for the white-light generation, because of the high sensitivity of the human eye in the green region of the spectrum. If high non-radiative losses are present, the recycling process can be enhanced by introducing more red and green QWs in the structure and by building a Bragg reflector for the blue wavelength. A possible way to enhance the external efficiencies of all three colours is to place the active regions into coupled microcavities.

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- [1] S. Nakamura and G. Fasol, *The Blue Laser Diode*, Springer, New York 1997.
 - [2] M. Asada, A. Kameyama, and Y. Suematsu, *IEEE J. Quantum Electron.*, **QE-20**, 745 (1998).
 - [3] M. Born and E. Wolf, *Principles of Optics*, Pergamon, Oxford 1970.
 - [4] H. Haug and S.W. Koch, *Quantum Theory of the Optical and Electronic Properties of Semiconductors*, World Scientific, Singapore, 1990.
 - [5] H. Benisty, R. Stanley and M. Mayer, *J. Opt. Soc. Am. A*, **15**, 1192 (1998).

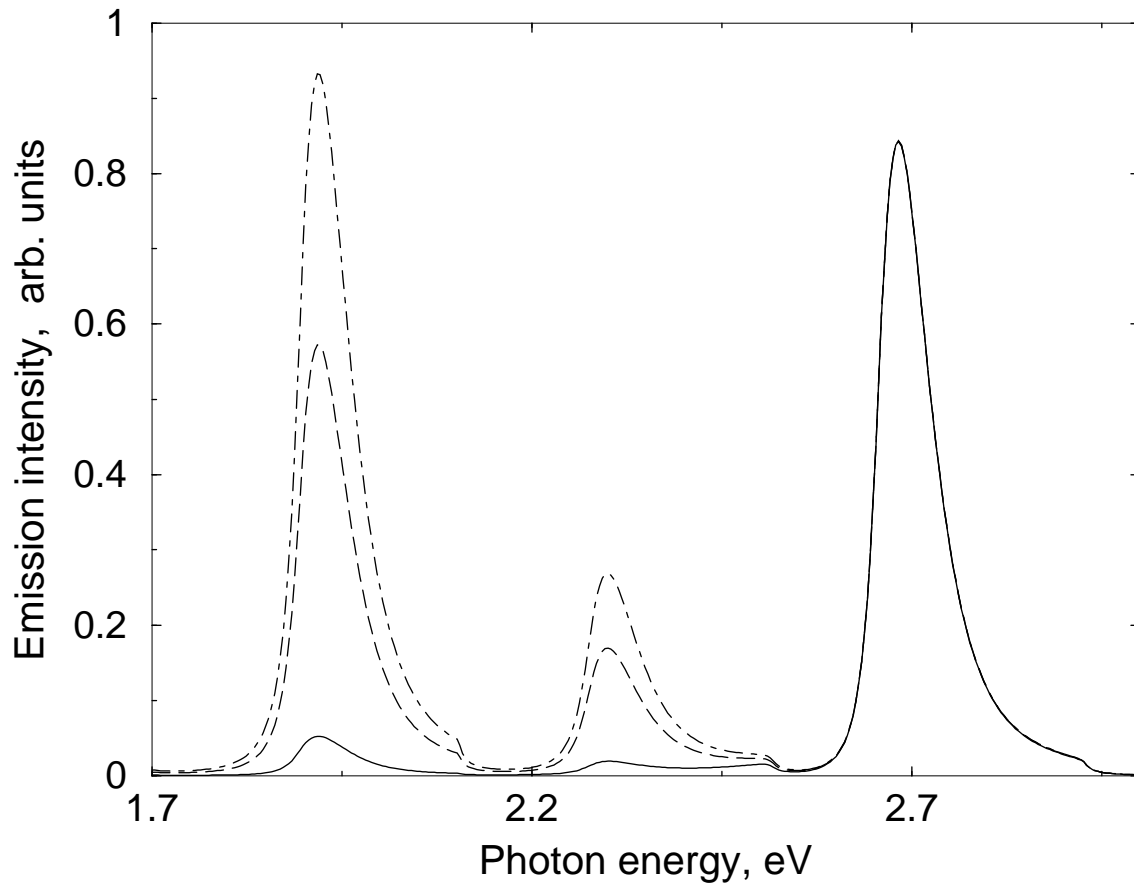


FIG. 1. Spontaneous emission spectra from a microcavity, containing three quantum wells. The non-radiative recombination times are 1 ns (solid line), 5 ns (dashed line) and 10 ns (dot-dashed line). The carrier density in the blue QW is the same for all three curves.